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(71) Applicant (for all designated States except US): **DEY, L.P.**
[US/US]; 2751 Napa Valley Corporate Drive, Napa, CA 94558 (US).

(72) Inventors; and

(75) Inventors/Applicants (for US only): **BANERJEE, Partha, S.** [IN/US]; 108 Rosario Court, San Ramon, CA 94583 (US). **CHAUDRY, Imitiaz, A.** [US/US]; 76 Golden Gate Circle, Napa, CA 94558 (US).

(74) Agents: **SEIDMAN, Stephanie, L.** et al.; Heller Ehrman White & McAuliffe LLP, 4350 La Jolla Village Drive, 6th Floor, San Diego, CA 92122-1246 (US).

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(54) Title: **FORMOTEROL/STEROID BRONCHODILATING COMPOSITIONS AND METHODS OF USE THEREOF**

(57) Abstract: Bronchodilating compositions and methods are provided. The compositions are intended for administration as a nebulized aerosol. In certain embodiments, the compositions contain formoterol, or a derivative thereof, and a steroidal anti-inflammatory agent. Methods for treatment, prevention, or amelioration of one or more symptoms of bronchoconstrictive disorders using the compositions provided herein are also provided.



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FORMOTEROL/STEROID BRONCHODILATING COMPOSITIONS AND METHODS OF USE THEREOF

RELATED APPLICATIONS

Benefit of priority is claimed to U.S. provisional patent application serial No. 60/284,607, filed April 17, 2001, to Banerjee *et al.*, and to U.S. application serial No. 09/887,496, filed June 22, 2001, to Banerjee *et al.* Where appropriate, the disclosures of the above-referenced applications are incorporated herein by reference in their entirety.

FIELD OF THE INVENTION

Compositions and methods are provided relating to treatment, prevention, or amelioration of one or more symptoms of bronchoconstrictive disorders. In particular, the compositions and methods herein include formoterol, and/or a derivative thereof, and a steroid, and/or a derivative thereof. The compositions are propellant-free, sterile unit dose or multidose inhalation solutions intended for administration via nebulization.

BACKGROUND OF THE INVENTION

Bronchoconstrictive disorders affect millions worldwide. Such disorders include asthma (including bronchial asthma, allergic asthma and intrinsic asthma, e.g., late asthma and airway hyper-responsiveness), chronic bronchitis and other chronic obstructive pulmonary diseases. Compounds having β_2 -adrenoreceptor agonist activity have been developed to treat these conditions. Such compounds include, but are not limited to, Albuterol (α^1 -(((1,1-dimethylethyl)amino)methyl)-4-hydroxy-1,3-benzenedimethanol); Bambuterol (dimethylcarbamic acid 5-(2-((1,1-dimethylethyl)amino)-1-hydroxyethyl)-1,3-phenylene ester); Bitolterol (4-methylbenzoic acid 4-(2-((1,1-dimethylethyl)amino)-1-hydroxyethyl)-1,2-phenylene ester); Broxaterol (3-bromo- α -(((1,1-dimethylethyl)amino)-methyl)-5-isoxazolemethanol); Isoproterenol (4-(1-hydroxy-2-((1-methyl-

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ethyl)amino)ethyl)-1,2-benzenediol); Trimetoquinol (1,2,3,4-tetrahydro-1-
((3,4,5-trimethoxyphenyl)methyl)-6,7-isoquinolinediol); Clenbuterol (4-
amino-3,5-dichloro- α -(((1,1-dimethylethyl)amino)methyl)benzenemethan-
ol); Fenoterol (5-(1-hydroxy-2-((2-(4-hydroxyphenyl)-1-methylethyl)-
5 amino)ethyl)-1,3-benzenediol); Formoterol (2-hydroxy-5-((1RS)-1-hydroxy-
2-(((1RS)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)formanilide);
(R,R)-Formoterol; Desformoterol ((R,R) or (S,S)-3-amino-4-hydroxy- α -((2-
(4-methoxyphenyl)-1-methylethyl)amino)methyl)benzenemethanol); Hexo-
prenaline (4,4'-(1,6-hexanediyl)-bis(imino(1-hydroxy-2,1-ethanediyl)))bis-
10 1,2-benzenediol); Isoetharine (4-(1-hydroxy-2-((1-methylethyl)amino)-
butyl)-1,2-benzenediol); Isoprenaline (4-(1-hydroxy-2-((1-methylethyl)-
amino)ethyl)-1,2-benzenediol); Metaproterenol (5-(1-hydroxy-2-((1-methyl-
ethyl)amino)ethyl)-1,3-benzenediol); Picumeterol (4-amino-3,5-dichloro- α -
(((6-(2-(2-pyridinyl)ethoxy)hexyl)amino)methyl)benzenemethanol); Pirbut-
15 erol (α^6 -(((1,1-dimethylethyl)amino)methyl)-3-hydroxy-2,6-pyridine-
methanol); Procaterol (((R*,S*)-(\pm))-8-hydroxy-5-(1-hydroxy-2-((1-
methylethyl)amino)butyl)-2(1H)-quinolinone); Reproterol ((7-(3-((2-(3,5-
dihydroxyphenyl)-2-hydroxyethyl)amino)propyl)-3,7-dihydro-1,3-dimethyl-
1H-purine-2,6-dione); Rimiterol (4-(hydroxy-2-piperidinylmethyl)-1,2-
20 benzenediol); Salbutamol ((\pm)- α^1 -(((1,1-dimethylethyl)amino)methyl)-4-
hydroxy-1,3-benzenedimethanol); (R)-Salbutamol; Salmeterol ((\pm)-4-
hydroxy- α^1 -(((6-(4-phenylbutoxy)hexyl)amino)methyl)-1,3-benzenedi-
methanol); (R)-Salmeterol; Terbutaline (5-(2-((1,1-dimethylethyl)amino)-1-
hydroxyethyl)-1,3-benzenediol); Tulobuterol (2-chloro- α -(((1,1-dimethyl-
25 ethyl)amino)methyl)benzenemethanol); and TA-2005 (8-hydroxy-5-((1R)-
1-hydroxy-2-(N-((1R)-2-(4-methoxyphenyl)-1-methylethyl)amino)ethyl)-
carbostyryl hydrochloride).

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These compounds are typically formulated for inhalation therapy. Aqueous or liquid formulations are preferred over formulations of solids. Powdered formulations are more difficult to administer, particularly to the young and elderly who are most often the patients in need of such therapy. Compounds, such as formoterol, which has many desirable properties, are not adequately stable in aqueous solutions to be formulated as liquids. Hence there is a need for formulations of compounds, such as formoterol, in a form that can be conveniently administered.

Other prophylactic therapeutics for use in treatment of bronchoconstrictive disorders include steroidal anti-inflammatory agents such as beclomethasone dipropionate (BDP), beclomethasone monopropionate (BMP), flunisolide, triamcenolone acetonide, dexamethasone, tipredane, ciclesonid, mometasone, mometasone furoate (Asmanex® Twisthaler™, Shering-Plough Corporation, Kenilworth, NJ), RPR 106541, fluticasone, fluticasone propionate and budesonide. These agents can be formulated for inhalation therapy.

Effective treatment of asthma and other bronchoconstrictive disorders often requires combination therapy. It is advantageous to administer combinations of bronchodilators and other agents, such as anti-steroidal agents. Since some the β_2 -adrenoreceptor agonist compounds are not available as aqueous or liquid formulations, combinations thereof with other agents, are not available as aqueous or liquid formulations. Since aqueous or liquid formulations are preferred, there is need to develop liquid formulations. Therefore, it is an object herein to provide stable liquid formulations of β_2 -adrenoreceptor agonist compounds. It is a further object herein to improve the stability of existing liquid formulations.

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SUMMARY OF THE INVENTION

Compositions and methods for treatment, prevention, or amelioration of one or more symptoms of bronchoconstrictive disorders are provided. The compositions provided herein are stable solutions of a
5 bronchodilating agent, and/or a derivative thereof, and a prophylactic therapeutic agent, including, but not limited to, a steroid and/or a derivative thereof. The compositions are formulated in a pharmacologically suitable fluid that contains water and that are stable during long term storage. The compositions are suitable for direct administration to a
10 subject in need thereof. Pharmacologically suitable fluids include, but are not limited to, polar fluids, including protic fluids. In certain embodiments herein, the compositions are aqueous solutions.

Compositions provided herein possess an estimated shelf-life of greater than 1, 2 or 3 months usage time at 25 °C and greater than or
15 equal to 1, 2 or 3 years storage time at 5 °C. In certain of these embodiments, using Arrhenius kinetics, >80% or >85% or >90% or >95% estimated bronchodilating agent remains after such storage. These compositions are particularly useful for administration via nebulization. In certain embodiments herein, the subject is a mammal. In
20 other embodiments, the subject is a human.

The compositions provided herein are formulated to remain stable over a relatively long period of time. For example, the compositions provided herein are stored between -15 °C and 25 °C, or between 2 °C and 8 °C. In one embodiment, the compositions are stored at 5 °C.

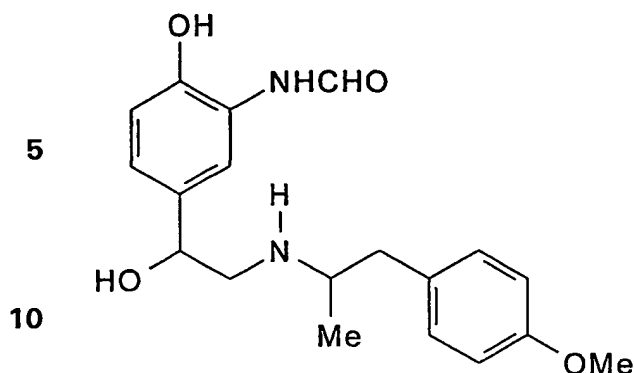
25 Among the bronchodilating agents for use herein are Albuterol (α^1 -(((1,1-dimethylethyl)amino)methyl)-4-hydroxy-1,3-benzenedimethanol); Bambuterol (dimethylcarbamic acid 5-(2-((1,1-dimethylethyl)amino)-1-hydroxyethyl)-1,3-phenylene ester); Bitolterol (4-methylbenzoic acid 4-(2-((1,1-dimethylethyl)amino)-1-hydroxyethyl)-1,2-phenylene ester); Broxa-

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terol (3-bromo- α -(((1,1-dimethylethyl)amino)methyl)-5-isoxazolemethanol); Isoproterenol (4-(1-hydroxy-2-((1-methylethyl)amino)ethyl)-1,2-benzenediol); Trimetoquinol (1,2,3,4-tetrahydro-1-((3,4,5-trimethoxyphenyl)-methyl)-6,7-isoquinolinediol); Clenbuterol (4-amino-3,5-dichloro- α -(((1,1-dimethylethyl)amino)methyl)benzenemethanol); Fenoterol (5-(1-hydroxy-2-((2-(4-hydroxyphenyl)-1-methylethyl)amino)ethyl)-1,3-benzenediol); Formoterol (2-hydroxy-5-((1R)-1-hydroxy-2-(((1R)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)formanilide); (R,R)-Formoterol; Desformoterol ((R,R) or (S,S)-3-amino-4-hydroxy- α -(((2-(4-methoxyphenyl)-1-methylethyl)amino)methyl)benzenemethanol); Hexoprenaline (4,4'-(1,6-hexanediyl)-bis(imino(1-hydroxy-2,1-ethanediyl)))bis-1,2-benzenediol); Isoetharine (4-(1-hydroxy-2-((1-methylethyl)amino)butyl)-1,2-benzenediol); Isoprenaline (4-(1-hydroxy-2-((1-methylethyl)amino)ethyl)-1,2-benzenediol); Metaproterenol (5-(1-hydroxy-2-((1-methylethyl)amino)ethyl)-1,3-benzenediol); Picumeterol (4-amino-3,5-dichloro- α -(((6-(2-(2-pyridinyl)ethoxy)hexyl)-amino)methyl)benzenemethanol); Pirbuterol (α^6 -(((1,1-dimethylethyl)-amino)methyl)-3-hydroxy-2,6-pyridinemethanol); Procaterol (((R^{*},S^{*})-(±)-8-hydroxy-5-(1-hydroxy-2-((1-methylethyl)amino)butyl)-2(1H)-quinolinone); Reproterol ((7-(3-((2-(3,5-dihydroxyphenyl)-2-hydroxyethyl)amino)-propyl)-3,7-dihydro-1,3-dimethyl-1H-purine-2,6-dione); Rimiterol (4-(hydroxy-2-piperidinylmethyl)-1,2-benzenediol); Salbutamol ((±)- α^1 -(((1,1-dimethylethyl)amino)methyl)-4-hydroxy-1,3-benzenedimethanol); (R)-Salbutamol; Salmeterol ((±)-4-hydroxy- α^1 -(((6-(4-phenylbutoxy)hexyl)-amino)methyl)-1,3-benzenedimethanol); (R)-Salmeterol; Terbutaline (5-(2-(((1,1-dimethylethyl)amino)-1-hydroxyethyl)-1,3-benzenediol); Tulobuterol (2-chloro- α -(((1,1-dimethylethyl)amino)methyl)benzenemethanol); and TA-2005 (8-hydroxy-5-((1R)-1-hydroxy-2-(N-((1R)-2-(4-methoxyphenyl)-1-methylethyl)amino)ethyl)carbostyryl hydrochloride).

Of particular interest herein is formoterol, having the formula:

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Formoterol for use in the compositions and methods provided

15 herein includes 2-hydroxy-5-((1RS)-1-hydroxy-2-(((1RS)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)formanilide; or a stereoisomer thereof; and also includes the single enantiomers 2-hydroxy-5-((1S)-1-hydroxy-2-(((1S)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)formanilide and 2-hydroxy-5-((1R)-1-hydroxy-2-(((1R)-2-(p-methoxyphenyl)-1-methylethyl)-

20 amino)ethyl)formanilide.

Prophylactic therapeutics for use in the compositions and methods herein include steroidal anti-inflammatory agents, including, but not limited to, beclomethasone dipropionate (BDP), beclomethasone monopropionate (BMP), flunisolide, triamcinolone acetonide,

25 dexamethasone, tipredane, ciclesonid, rofleponide, mometasone, mometasone furoate (Asmanex® Twisthaler™, Shering-Plough Corporation, Kenilworth, NJ), RPR 106541, fluticasone or fluticasone propionate and budesonide, or derivatives thereof. In one embodiment, the steroidal anti-inflammatory is fluticasone, fluticasone propionate, budesonide, or a

30 derivative thereof.

In certain embodiments, the compositions are administered via nebulization. Administration of a nebulized aerosol is preferred over the

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use of dry powders for inhalation in certain subject populations, including pediatric and geriatric groups.

- In one embodiment, the compositions for use in the methods provided herein contain a pharmaceutically acceptable derivative of
- 5 formoterol. In another embodiment, the compositions for use in the methods provided herein contain a pharmaceutically acceptable salt of formoterol. Pharmaceutically acceptable salts include, but are not limited to, salts of mineral acids, such as but not limited to hydrochlorides and sulfates; and salts of organic acids, such as but not limited to acetates,
- 10 lactates, malates, tartrates, citrates, ascorbates, succinates, butyrates, valerates and fumarates. In one embodiment, the compositions for use in the methods provided herein contain formoterol fumarate or formoterol fumarate dihydrate. In another embodiment, the compositions for use in the methods provided herein contain formoterol tartrate.
- 15 In another embodiment, the compositions for use in the methods provided herein contain budesonide or fluticasone propionate. Compositions containing formoterol, budesonide and a fluoroalkane propellant are disclosed in U.S. Patent No. 6,004,537. Compositions containing formoterol and budesonide for dry powder inhalation or
- 20 metered dose inhalation are disclosed in U.S. Patent Nos. 5,674,860 and 5,972,919. These references do not disclose the compositions provided herein that are formulated in a pharmacologically suitable fluid, as defined herein, that contains water and that are stable during long term storage.

- Also provided herein are combinations containing compositions
- 25 provided herein. The compositions may be formulated separately or mixed in a single composition. The compositions contained in the combinations may be administered sequentially or intermittently. The compositions contained in the combinations can be mixed prior to use or can be formulated as a single composition. The combinations may

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further include a nebulizer. The combinations can be packaged as kits, which optionally contain other components, including instructions for use of the nebulizer and/or instructions for mixing the compositions if provided separately.

- 5 Any nebulizer is contemplated for use in the kits and methods provided herein. In particular, the nebulizers for use herein nebulize liquid formulations, including the compositions provided herein, containing no propellant. The nebulizer may produce the nebulized mist by any method known to those of skill in the art, including, but not limited to,
- 10 compressed air, ultrasonic waves, or vibration. The nebulizer may further have an internal baffle. The internal baffle, together with the housing of the nebulizer, selectively removes large droplets from the mist by impaction and allows the droplets to return to the reservoir. The fine aerosol droplets thus produced are entrained into the lung by the inhaling
- 15 air/oxygen.

- Methods for the treatment, prevention, or amelioration of one or more symptoms of bronchoconstrictive disorders, including, but not limited to, asthma, including, but not limited to, bronchial asthma, allergic asthma and intrinsic asthma, e.g., late asthma and airway hyper-
- 20 responsiveness; chronic bronchitis; and other chronic obstructive pulmonary diseases are provided. The methods involve administering an effective amount of a pharmaceutical composition provided herein to a subject in need of such treatment.

- Articles of manufacture, containing packaging material, a
- 25 composition provided herein, which is useful for treatment, prevention or amelioration of one or more symptoms of diseases or disorders associated with undesired and/or uncontrolled bronchoconstriction, and a label that indicates that the composition is used for treatment, prevention or amelioration of one or more symptoms of diseases or disorders associated

with undesired and/or uncontrolled bronchoconstriction, are also provided.

DETAILED DESCRIPTION OF PREFERRED EMBODIMENTS

Definitions

- 5 Unless defined otherwise, all technical and scientific terms used herein have the same meaning as is commonly understood by one of ordinary skill in the art to which this invention belongs. All patents, applications, published applications and other publications are incorporated by reference in their entirety. In the event that there are a
- 10 plurality of definitions for a term herein, those in this section prevail unless stated otherwise.

- As used herein, formoterol refers to 2-hydroxy-5-((1RS)-1-hydroxy-2-(((1RS)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)formanilide; or a stereoisomer thereof. The term formoterol also refers to the single
- 15 enantiomers 2-hydroxy-5-((1S)-1-hydroxy-2-(((1S)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)formanilide and 2-hydroxy-5-((1R)-1-hydroxy-2-(((1R)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)formanilide.

- As used herein, formoterol fumarate refers to a salt of formoterol having the formula (formoterol)*½ fumarate. Formoterol fumarate
- 20 dihydrate refers to the dihydrate of formoterol fumarate.

- As used herein, formoterol free base refers to the neutral, anhydrous form of formoterol. Thus, a recitation that a composition contains, e.g., 59 µg/mL of formoterol free base means that the composition contains 59 µg/mL of neutral, anhydrous formoterol. Such
- 25 compositions may be prepared using a derivative of formoterol.

As used herein, budesonide refers to (RS)-11β,16α,17,21-tetrahydroxypregna-1,4-diene-3,20-dione cyclic 16,17-acetal with butraldehyde. The term budesonide refers to the (R) isomer, the (S) isomer, and mixtures thereof.

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As used herein, fluticasone refers to (6 α ,9 α ,11 β ,16 α ,17 α)-6,9-difluoro-11-hydroxy-16-methyl-3-oxo-17-hydroxyandrost-1,4-diene-17-carbothioic acid, S-fluoromethyl ester. Fluticasone propionate refers to (6 α ,9 α ,11 β ,16 α ,17 α)-6,9-difluoro-11-hydroxy-16-methyl-3-oxo-17-(1-oxo-
5 propoxy)androst-1,4-diene-17-carbothioic acid, S-fluoromethyl ester.

As used herein, an aerosol is liquid or particulate matter dispersed in air. Aerosols include dispersions of liquids, including aqueous and other solutions, and solids, including powders, in air.

As used herein, a nebulized solution refers to a solution that is
10 dispersed in air to form an aerosol. Thus, a nebulized solution is a particular form of an aerosol.

As used herein, a nebulizer is an instrument that is capable of generating very fine liquid droplet for inhalation into the lung. Within this instrument, the nebulizing liquid or solution is atomized into a mist of
15 droplets with a broad size distribution by methods known to those of skill in the art, including, but not limited to, compressed air, ultrasonic waves, or a vibrating orifice. Nebulizers may further contain, e.g., a baffle which, along with the housing of the instrument, selectively removes large droplets from the mist by impaction. Thus, the mist inhaled into the lung
20 contains fine aerosol droplets.

As used herein, a pharmacologically suitable fluid is a solvent suitable for pharmaceutical use which is not a liquified propellant gas. Exemplary pharmacologically suitable fluids include polar fluids, including protic fluids such as water.

25 As used herein, a combination refers to any association between two or among more items.

As used herein, fluid refers to any composition that can flow. Fluids thus encompass compositions that are in the form of semi-solids,

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pastes, solutions, aqueous mixtures, gels, lotions, creams and other such compositions.

As used herein, a mixture is a mutual incorporation of two or more substances, without chemical union, the physical characteristics of each
5 of the components being retained.

As used herein, the stability of a composition provided herein refers to the length of time at a given temperature that greater than 80%, 85%, 90% or 95% of the initial amount of active ingredient, e.g., formoterol, is present in the composition. Thus, for example, a composition that is
10 stable for 30 days at 25 °C would have greater than 80%, 85%, 90% or 95% of the initial amount of active ingredient present in the composition at 30 days following storage at 25 °C.

As used herein, pharmaceutically acceptable derivatives of a compound include salts, esters, enol ethers, enol esters, acids, bases,
15 solvates, hydrates or prodrugs thereof. Such derivatives may be readily prepared by those of skill in this art using known methods for such derivatization. The compounds produced may be administered to animals or humans without substantial toxic effects and either are pharmaceutically active or are prodrugs. Pharmaceutically acceptable
20 salts include, but are not limited to, amine salts, such as but not limited to N,N'-dibenzylethylenediamine, chlorprocaine, choline, ammonia, diethanolamine and other hydroxyalkylamines, ethylenediamine, N-methylglucamine, procaine, N-benzylphenethylamine, 1-para-chlorobenzyl-2-pyrrolidin-1'-ylmethylbenzimidazole, diethylamine and other alkylamines,
25 piperazine and tris(hydroxymethyl)aminomethane; alkali metal salts, such as but not limited to lithium, potassium and sodium; alkali earth metal salts, such as but not limited to barium, calcium and magnesium; transition metal salts, such as but not limited to zinc; and other metal salts, such as but not limited to sodium hydrogen phosphate and

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disodium phosphate; and also including, but not limited to, salts of mineral acids, such as but not limited to hydrochlorides and sulfates; and salts of organic acids, such as but not limited to acetates, lactates, malates, tartrates, citrates, ascorbates, succinates, butyrates, valerates
5 and fumarates. Pharmaceutically acceptable esters include, but are not limited to, alkyl, alkenyl, alkynyl, aryl, heteroaryl, aralkyl, heteroaralkyl, cycloalkyl and heterocyclyl esters of acidic groups, including, but not limited to, carboxylic acids, phosphoric acids, phosphinic acids, sulfonic acids, sulfinic acids and boronic acids. Pharmaceutically acceptable enol
10 ethers include, but are not limited to, derivatives of formula $C=C(OR)$ where R is hydrogen, alkyl, alkenyl, alkynyl, aryl, heteroaryl, aralkyl, heteroaralkyl, cycloalkyl or heterocyclyl. Pharmaceutically acceptable enol esters include, but are not limited to, derivatives of formula $C=C(OC(O)R)$ where R is hydrogen, alkyl, alkenyl, alkynyl, aryl,
15 heteroaryl, aralkyl, heteroaralkyl, cycloalkyl or heterocyclyl. Pharmaceutically acceptable solvates and hydrates are complexes of a compound with one or more solvent or water molecule, preferably 1 to about 100, more preferably 1 to about 10, most preferably one to about 2, 3 or 4, solvent or water molecules. Formoterol salts and hydrates are
20 used in certain embodiments herein.

As used herein, treatment means any manner in which one or more of the symptoms of a condition, disorder or disease are ameliorated or otherwise beneficially altered. Treatment also encompasses any pharmaceutical use of the compositions herein, such as use for treating
25 cancer.

As used herein, amelioration of the symptoms of a particular disorder by administration of a particular pharmaceutical composition refers to any lessening, whether permanent or temporary, lasting or

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transient that can be attributed to or associated with administration of the composition.

As used herein, a prodrug is a compound that, upon in vivo administration, is metabolized or otherwise converted to the biologically, pharmaceutically or therapeutically active form of the compound. To produce a prodrug, the pharmaceutically active compound is modified such that the active compound will be regenerated by metabolic processes. The prodrug may be designed to alter the metabolic stability or the transport characteristics of a drug, to mask side effects or toxicity, to improve the flavor of a drug or to alter other characteristics or properties of a drug. By virtue of knowledge of pharmacodynamic processes and drug metabolism in vivo, those of skill in this art, once a pharmaceutically active compound is known, can design prodrugs of the compound (see, e.g., Nogrady (1985) Medicinal Chemistry A Biochemical Approach, Oxford University Press, New York, pages 388-392).

It is to be understood that the compounds for use in the compositions and methods provided herein may contain chiral centers. Such chiral centers may be of either the (R) or (S) configuration, or may be a mixture thereof. Thus, the compounds for use in the compositions provided herein may be enantiomerically pure, or be stereoisomeric or diastereomeric mixtures. It is to be understood that the chiral centers of the compounds provided herein may undergo epimerization in vivo. Thus, one of skill in the art will recognize that administration of a compound in its (R) form is equivalent, for compounds that undergo epimerization in vivo, to administration of the compound in its (S) form.

As used herein, bronchoconstriction refers to a reduction in the caliber of a bronchus or bronchi.

As used herein, undesired and/or uncontrolled bronchoconstriction refers to bronchoconstriction that results in or from a pathological

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symptom or condition. Pathological conditions include, but are not limited to, asthma and chronic obstructive pulmonary disease (COPD). Pathological symptoms include, but are not limited to, asthma and COPD.

As used herein, the statement that a composition is stable during
5 "long term storage" means that the composition is suitable for administration to a subject in need thereof when it has an estimated shelf-life of greater than 1, 2 or 3 months usage time at 25 °C and greater than or equal to 1, 2 or 3 years storage time at 5 °C. In certain embodiments herein, using Arrhenius kinetics, >80% or >85% or
10 >90% or >95% estimated bronchodilating agent remains after such storage.

A. Formoterol, Budesonide and Fluticasone Propionate

1. Formoterol

Formoterol (2-hydroxy-5-((1RS)-1-hydroxy-2-(((1RS)-2-(p-methoxy-
15 phenyl)-1-methylethyl)amino)ethyl)formanilide) is derived from adrenaline and, as noted above, is used as a β_2 -stimulator in inhalation therapy of respiratory diseases, particularly for the treatment of bronchial asthma. It has been reported that in patients with reversible obstructive respiratory diseases, formoterol has a bronchodilatory effect. This effect has a
20 relatively rapid onset (approximately 1-3 minutes) and a relatively long duration (greater than 12 hours). Formoterol inhibits the release of leukotrienes and other messenger substances involved with inflammation, such as histamines. In addition, formoterol may bring about a hyperglycaemic activity.

25 To date, formoterol has been formulated as a dry powder and administered via devices such as the Turbuhaler® and the Aerolizer®. See, e.g., Seberova *et al.* (2000) *Respir. Med.* 94(6):607-611; Lotvall *et al.* (1999) *Can. Respir. J.* 6(5):412-416; Campbell *et al.* (1999) *Respir. Med.* 93(4):236-244; Nightingale *et al.* (1999) *Am. J. Respir. Crit. Care*

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Med. 159(6):1786-1790; Lecaillon et al. (1999) Eur. J. Clin. Pharmacol. 55(2):131-138; Bartow et al. (1998) Drugs 55(2):303-322; Ekstrom et al. (1998) Respir. Med. 92(8):1040-1045; Ringdal et al. (1998) Respir. Med. 92(8):1017-1021; Totterman et al. (1998) Eur. Respir. J. 12(3):573-579; Palmqvist et al. (1997) Eur. Respir. J. 10(11):2484-2489; Nielsen et al. (1997) Eur. Respir. J. 10(9):2105-2109; Ullman et al. (1996) Allergy 51(10):745-748; Selroos et al. (1996) Clin. Immunother. 6:273-299; and Schreurs et al. (1996) Eur. Respir. J. 9(8):1678-1683.

- 10** Formoterol is also available as a tablet and a dry syrup in certain areas of the world (e.g., Atock®, marcketed by Yamanouchi Pharmaceutical Co. Ltd., Japan). Formoterol formulations are also available in other areas (e.g., Europe and U.S.) for propellant-based metered dose inhalers and dry powder inhalers (e.g., Turbuhaler®,
- 15** Aerolizer® and Foradil Aerolizer®). None of these formulations are water based. Sterile, stable, aqueous based inhalation solutions of formoterol for nebulization are not available, nor have they been reported.

Compositions containing formoterol in combination with other active ingredients have been disclosed. See, e.g., 5,668,110,

- 20** 5,683,983, 5,677,280 and 5,654,276 (formoterol and IL-5 inhibitors), 6,136,603 (formoterol and antisense modulators of IL-5), 5,602,110 (formoterol and milrinone), 5,525,623 (formoterol and a tryptase inhibitor), 5,691,336, 5,877,191, 5,929,094, 5,750,549 and 5,780,467 (formoterol and a tachykinin receptor antagonist); and International Patent
- 25** Application Publication Nos. WO 99/00134 (formoterol and rofleponide) and WO 99/36095 (formoterol and a dopamine D₂ receptor agonist).

Other compositions containing formoterol have been disclosed in U.S. Patent Nos. 5,677,809, 6,126,919, 5,733,526, 6,071,971,

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6,068,833, 5,795,564, 6,040,344, 6,041,777, 5,874,481, 5,965,622 and 6,161,536.

U.S. Patent No. 6,150,418 discloses a "liquid active substance concentrate" containing formoterol in the form of its free base or in the
5 form of one of the pharmacologically acceptable salts or addition products (adducts) thereof as active substance. This "liquid active substance concentrate" is reported to be a concentrated (i.e., greater than 10 mg/mL, preferably 75 to 500 mg/mL) solution or suspension that is stable for a period of several months possibly up to several years
10 without any deterioration in the pharmaceutical quality. This patent teaches that it is the high concentration that allows for the stability of the concentrate. The "liquid active substance concentrate" is not suitable for direct administration to a subject.

U.S. Patent No. 6,040,344 discloses an aqueous aerosol formula-
15 tion of formoterol tartrate for use in a nebulizer. This patent states that the formulation disclosed therein is not attractive for long term storage.

2. Budesonide and Fluticasone Propionate

Budesonide, (RS)-11 β ,16 α ,17,21-tetrahydroxypregna-1,4-diene-3,20-dione cyclic 16,17-acetal with butraldehyde, is an anti-
20 inflammatory, synthetic corticosteroid. Fluticasone propionate, (6 α ,11 β ,16 α ,17 α)-6,9,-difluoro-11-hydroxy-16-methyl-3-oxo-17-(1-oxopropoxy)androsta-1,4-diene-17-carbothioic acid, S-fluoromethyl ester, is a synthetic fluorinated corticosteroid, and is known for topical dermatologic use. Topical corticosteroids constitute a class of primarily
25 synthetic steroids used as anti-inflammatory and antipruritic agents. The mechanism of the anti-inflammatory activity of topical steroids, in general, is unclear. However, corticosteroids are thought to act by the induction of phospholipase A2 inhibitory proteins (lipocortins). It is

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postulated that these proteins control the biosynthesis of prostaglandins and leukotrienes by inhibiting the release of arachidonic acid.

B. Compositions for use in treatment, prevention, or amelioration of one or more symptoms of bronchoconstrictive disorders

- 5 Pharmaceutical compositions containing a β_2 -adrenoreceptor agonist and a steroid for administration via nebulization are provided. The compositions are sterile filtered and filled in vials, including unit dose vials providing sterile unit dose formulations which are used in a nebulizer and suitably nebulized. Each unit dose vial is sterile and is suitably nebulized
- 10 without contaminating other vials or the next dose. The bulk sterile formulation is sterilized by steam, gamma radiation or is prepared using sterile steroidal powder.

- The unit dose vials are formed in a form-fill-seal machine or by any other suitable method known to those of skill in the art. The vials may be
- 15 made of plastic materials that are suitably used in these processes. For example, plastic materials for preparing the unit dose vials include, but are not limited to, low density polyethylene, high density polyethylene, polypropylene and polyesters. In one embodiment, the plastic material is low density polyethylene.

- 20 In one embodiment, the β_2 -adrenoreceptor agonist is formoterol, or a pharmaceutically acceptable derivative thereof. In other embodiments, the formoterol for use in the compositions provided herein is formoterol fumarate. Formoterol refers to 2-hydroxy-5-((1RS)-1-hydroxy-2-(((1RS)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)formanilide; or a stereo-
- 25 isomer thereof. The term formoterol also refers herein to the single enantiomers 2-hydroxy-5-((1S)-1-hydroxy-2-(((1S)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)formanilide and 2-hydroxy-5-((1R)-1-hydroxy-2-(((1R)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)formanilide.

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In one embodiment, the compositions contain, in addition to a steroidal anti-inflammatory agent, including, but not limited to, budesonide and fluticasone propionate, formoterol free base at a concentration of about 5 $\mu\text{g/mL}$ to about 2 mg/mL . In other

5 embodiments, the maximum concentration of formoterol free base in the compositions is 1.5 mg/mL . In further embodiments, the concentration of formoterol free base in the compositions is about 10 $\mu\text{g/mL}$ to about 1 mg/mL , or about 50 $\mu\text{g/mL}$ to about 200 $\mu\text{g/mL}$. In other embodiments, the compositions contain formoterol fumarate at a concentration of about

10 80 $\mu\text{g/mL}$ up to about 175 to 200 $\mu\text{g/mL}$. In further embodiments, the compositions contain formoterol fumarate at a concentration of about 90 $\mu\text{g/mL}$ up to about 125 to 150 $\mu\text{g/mL}$. The formoterol fumarate is formulated, in certain compositions provided herein, at a concentration of about 100 $\mu\text{g/mL}$. The formoterol fumarate is formulated, in other

15 compositions provided herein, at a concentration of about 85 $\mu\text{g/mL}$ or about 170 $\mu\text{g/mL}$. In one embodiment, the formoterol fumarate is formulated for single dosage administration via nebulization at a concentration of about 100 $\mu\text{g/mL}$. In another embodiment, the compositions contain formoterol free base at a concentration of about 40

20 to about 150 $\mu\text{g/mL}$, particularly about 59 or about 118 $\mu\text{g/mL}$.

The compositions provided herein further contain, in addition to a β_2 -adrenoreceptor agonist, including formoterol, a steroidal anti-inflammatory agent, including, but not limited to, budesonide or fluticasone propionate. Budesonide is (RS)-11 β ,16 α ,17,21-

25 tetrahydroxypregna-1,4-diene-3,20-dione cyclic 16,17-acetal with butraldehyde. Budesonide also refers to the (R) isomer, the (S) isomer, and mixtures thereof. Fluticasone propionate refers to (6 α ,11 β ,16 α ,17 α)-6,9-difluoro-11-hydroxy-16-methyl-3-oxo-17-(1-oxopropoxy)androsta-1,4-diene-17-carbothioic acid, S-fluoromethyl ester.

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In one embodiment, the compositions contain budesonide or fluticasone propionate at a concentration of about 5 $\mu\text{g/mL}$ to about 2 mg/mL. In another embodiment, the compositions contain budesonide at a concentration of about 75 $\mu\text{g/mL}$ to about 500 $\mu\text{g/mL}$, or about 125 $\mu\text{g/mL}$ to about 500 $\mu\text{g/mL}$. In certain embodiments, the compositions contain budesonide at a concentration of about 125 $\mu\text{g/mL}$ or 250 $\mu\text{g/mL}$. In another embodiment, the compositions contain fluticasone propionate at a concentration of about 75 $\mu\text{g/mL}$ to about 1000 $\mu\text{g/mL}$, or about 250 $\mu\text{g/mL}$ to about 1000 $\mu\text{g/mL}$. In further embodiments, the compositions contain fluticasone propionate at a concentration of about 125 $\mu\text{g/mL}$ or about 250 $\mu\text{g/mL}$.

The compositions are formulated as solutions or suspensions with a pharmacologically suitable fluid. Pharmacologically suitable fluids include, but are not limited to, polar solvents, including, but not limited to, compounds that contain hydroxyl groups or other polar groups. Such solvents include, but are not limited to, water or alcohols, such as ethanol, isopropanol, and glycols including propylene glycol, polyethylene glycol, polypropylene glycol, glycol ether, glycerol and polyoxyethylene alcohols.

Polar solvents also include protic solvents, including, but not limited to, water, aqueous saline solutions with one or more pharmaceutically acceptable salt(s), alcohols, glycols or a mixture thereof. For a saline solution as the solvent or as a component thereof, particularly suitable salts are those which display no or only negligible pharmacological activity after administration.

In the embodiments herein, the compositions have a pH of about 2.0 to about 8.0. The particular pH of a given composition for long term storage provided herein may be determined empirically using standard stability assays well known to those of skill in the art (see, e.g., the

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Examples). For example, in certain embodiments, the compositions have a pH of about 4.0 to about 6.0, or about 4.5 to about 5.5. In certain of the above embodiments, the compositions are formulated at a pH of about 4, 4.4 or 4.6 up to about 5.5, 5.7 or 6. In other embodiments, the pH is about 5.0. It has been found herein that the rate constant for decomposition of an aqueous solution of formoterol is dependent on pH. The rate constant (k_{obs}) at 60 °C at a pH of 3, 4, 5 and 7 is approximately 0.62, 0.11, 0.044 and 0.55 day⁻¹, respectively. Therefore, the decomposition of formoterol in aqueous solution at 60 °C at a buffer concentration of 5 mM and an ionic strength of 0.05 is slowest at a pH of about 5.0.

The solubility of formoterol in aqueous solution has been found herein to be dependent on pH. Thus, at a pH of between about 5 and about 7, the aqueous solubility of formoterol at ambient temperature is approximately 2.2 mg/mL. At a pH of about 4, the aqueous solubility of formoterol at ambient temperature is approximately 3 mg/mL, while at a pH of about 3, the aqueous solubility of formoterol at ambient temperature is about 4.8 mg/mL. The solubility of formoterol in pure water, for example, high performance liquid chromatography (HPLC) water, at ambient temperature is approximately 2 mg/mL.

In other of the above embodiments, the compositions further contain a buffer, including, but not limited to, citric acid/phosphate, acetate, barbital, borate, Britton-Robinson, cacodylate, citrate, collidine, formate, maleate, McIlvaine, phosphate, Prideaux-Ward, succinate, citrate-phosphate-borate (Teorell-Stanhagen), veronal acetate, MES (2-(N-morpholino)ethanesulfonic acid), BIS-TRIS (bis(2-hydroxyethyl)iminotris-(hydroxymethyl)methane), ADA (N-(2-acetamido)-2-iminodiacetic acid), ACES (N-(carbamoylmethyl)-2-aminoethanesulfonic acid), PIPES (piperazine-N,N'-bis(2-ethanesulfonic acid)), MOPSO (3-(N-morpholino)-2-

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hydroxypropanesulfonic acid), BIS-TRIS PROPANE (1,3-bis(tris(hydroxymethyl)methylamino)propane), BES (N,N-bis(2-hydroxyethyl)-2-aminoethanesulfonic acid), MOPS (3-(N-morpholino)propanesulfonic acid), TES (N-tris(hydroxymethyl)methyl-2-aminoethanesulfonic acid), HEPES (N-(2-hydroxyethyl)piperazine-N'-(2-ethanesulfonic acid), DIPSO (3-(N,N-bis(2-hydroxyethyl)amino)-2-hydroxypropanesulfonic acid), MOBS (4-(N-morpholino)butanesulfonic acid), TAPSO (3-(N-tris(hydroxymethyl)methylamino)-2-hydroxypropanesulfonic acid), TRIZMA® (tris(hydroxymethyl)aminomethane), HEPPSO (N-(2-hydroxyethyl)piperazine-N'-(2-hydroxypropanesulfonic acid), POPSO (piperazine-N,N'-bis(2-hydroxypropanesulfonic acid)), TEA (triethanolamine), EPPS (N-(2-hydroxyethyl)piperazine-N'-(3-propanesulfonic acid), TRICINE (N-tris(hydroxymethyl)methylglycine), GLY-GLY (glycylglycine), BICINE (N,N-bis(2-hydroxyethyl)glycine), HEPBS (N-(2-hydroxyethyl)piperazine-N'-(4-butanefulfonic acid)), TAPS (N-tris(hydroxymethyl)methyl-3-amino-
15 propanesulfonic acid), AMPD (2-amino-2-methyl-1,3-propanediol), and/or any other buffers known to those of skill in the art. In one embodiment, the buffer is citric acid/phosphate buffer, acetate buffer, citrate buffer or phosphate buffer. In another embodiment, the buffer is a citrate buffer
20 (citric acid/sodium citrate). The buffer concentration has been found herein to affect the stability of the composition. Buffer concentrations for use herein include from about 0 or 0.01 mM to about 150 mM, or about 1 mM to about 20 mM. In one embodiment, the buffer concentration is about 5 mM. In another embodiment, the buffer
25 concentration is about 1 mM to about 50 mM, or about 20 mM. The kinetic-pH profile of formoterol is dependent on buffer concentration. At low and approximately neutral conditions, increasing the buffer concentration from 5 mM to 20 mM increased the rated constant of decomposition significantly. However, no noticeable differences in rate

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constant were observed in the pH region of about 4.5 to about 5.5 with increasing buffer concentration from 5 mM to 20 mM. The particular buffer and buffer concentration of a given composition for long term storage provided herein may be determined empirically using standard
5 stability assays well known to those of skill in the art (see, e.g., the Examples).

The ionic strength of the compositions provided herein also has been found herein to affect the stability of the composition. Ionic strengths of the compositions provided herein are from about 0 to about
10 0.4, or from about 0.05 to about 0.16. Compositions having a lower ionic strength exhibit improved stability over formulations having higher ionic strength. The rate constant of decomposition was essentially the same at ionic strength 0.05 to 0.1, but increased to some extent at ionic strength of 0.2. The particular ionic strength of a given composition for
15 long term storage provided herein may be determined empirically using standard stability assays well known to those of skill in the art (see, e.g., the Examples).

In embodiments where the pharmacologically suitable fluid is a saline solution, tonicity adjusting agents may be added to provide the
20 desired ionic strength. Tonicity adjusting agents for use herein include those which display no or only negligible pharmacological activity after administration. Both inorganic and organic tonicity adjusting agents may be used in the compositions provided herein. Tonicity adjusting agents include, but are not limited to, ammonium carbonate, ammonium chloride,
25 ammonium lactate, ammonium nitrate, ammonium phosphate, ammonium sulfate, ascorbic acid, bismuth sodium tartrate, boric acid, calcium chloride, calcium disodium edetate, calcium gluconate, calcium lactate, citric acid, dextrose, diethanolamine, dimethylsulfoxide, edetate disodium, edetate trisodium monohydrate, fluorescein sodium, fructose, galactose,

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- glycerin, lactic acid, lactose, magnesium chloride, magnesium sulfate, mannitol, polyethylene glycol, potassium acetate, potassium chlorate, potassium chloride, potassium iodide, potassium nitrate, potassium phosphate, potassium sulfate, propylene glycol, silver nitrate, sodium acetate, sodium bicarbonate, sodium biphosphate, sodium bisulfite, sodium borate, sodium bromide, sodium cacodylate, sodium carbonate, sodium chloride, sodium citrate, sodium iodide, sodium lactate, sodium metabisulfite, sodium nitrate, sodium nitrite, sodium phosphate, sodium propionate, sodium succinate, sodium sulfate, sodium sulfite, sodium tartrate, sodium thiosulfate, sorbitol, sucrose, tartaric acid, triethanolamine, urea, urethan, uridine and zinc sulfate. In certain embodiments, the tonicity adjusting agent is sodium chloride, which is present at a concentration of from about 0 mg/mL to about 10, 15 or 20 mg/mL. In further embodiments, the compositions contain sodium chloride at a concentration of from about 0 mg/mL to about 7.5 mg/mL. In another embodiment, the compositions contain sodium chloride at a concentration of 0 mg/mL, 1.5 mg/mL, 6.8 mg/mL or 7.5 mg/mL. In these embodiments, the pharmacologically suitable fluid is aqueous saline.
- 20 The storage temperature of the compositions provided herein also has been found herein to affect the stability of the composition. Compositions stored at a lower temperature exhibit improved stability over formulations stored at higher temperatures. The effect of temperature on the rate constant of decomposition at pH 5, a buffer concentration of 5 mM, and an ionic strength of 0.05, was linear according to Arrhenius kinetics, i.e., when $\ln k_{obs}$ was plotted against $1/T$, where T is the temperature in degree Kelvin.

The estimated shelf-life of formoterol in the compositions provided herein is significantly greater than that reported for known formoterol

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compositions. The estimated shelf-life of formoterol in the compositions provided herein is about 6.2 years at 5 °C and about 7.5 months at 25 °C. The estimated formoterol concentrations in the compositions provided herein as a function of storage time at 5 °C and usage time at 5 25 °C was determined. It is estimated that greater than 90% of the initial formoterol present in the composition remains after 3 months of usage time at 25 °C and 3 years of storage time at 5 °C as well as after 0.5 months of usage time at 25 °C and 1 year of storage time at 5 °C.

In certain embodiments, the compositions provided herein are 10 prepared containing formoterol fumarate at a nominal concentration of 0.1 mg/mL at the indicated pH and citric acid/phosphate buffer concentrations. The solutions were stored at 60 °C. In these compositions, formoterol is relatively more stable at a pH from about 4 to about 5, and is also more stable at lower buffer concentration.

15 The compositions provided herein also may include excipients and additives. The particular excipient or additive for use in the compositions for long term storage provided herein may be determined empirically using methods well known to those of skill in the art (see, e.g., the Examples). Excipients and additives are any pharmacologically suitable and 20 therapeutically useful substance which is not an active substance. Excipients and additives generally have no pharmacological activity, or at least no undesirable pharmacological activity. The excipients and additives include, but are not limited to, surfactants, stabilizers, complexing agents, antioxidants, or preservatives which prolong the 25 duration of use of the finished pharmaceutical formulation, flavorings, vitamins, or other additives known in the art. Complexing agents include, but are not limited to, ethylenediaminetetraacetic acid (EDTA) or a salt thereof, such as the disodium salt, citric acid, nitrilotriacetic acid and the salts thereof. In one embodiment, the complexing agent is EDTA. In

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another embodiment, particularly in the suspension formulations provided herein, the complexing agent is sodium edetate. In these embodiments, the compositions contain sodium edetate at a concentration of about 0.05 mg/mL to about 0.5 mg/mL, or about 0.1 mg/mL to about 0.2 mg/mL.

Preservatives include, but are not limited to, those that protect the solution from contamination with pathogenic particles, including benzalkonium chloride or benzoic acid, or benzoates such as sodium benzoate. Antioxidants include, but are not limited to, vitamins, provitamins, ascorbic acid, vitamin E or salts or esters thereof.

In certain embodiments herein, particularly in the solution formulations provided herein, the compositions contain vitamin E TPGS (d- α -tocopheryl polyethylene glycol 1000 succinate). In these embodiments, vitamin E TPGS is present at a concentration of about 0 mg/mL to about 100 mg/mL, or about 5 mg/mL to about 50 mg/mL. In certain embodiments herein, the compositions contain vitamin E TPGS at a concentration of 10 mg/mL, 20 mg/mL, 30 mg/mL or 50 mg/mL.

The compositions provided herein also may include a cosolvent, which increases the solubility of additives or the active ingredient(s). The particular cosolvent for use in the compositions for long term storage provided herein may be determined empirically using methods well known to those of skill in the art (see, e.g., the Examples). Cosolvents for use herein include, but are not limited to, hydroxylated solvents or other polar solvents, such as alcohols such as isopropyl alcohol, glycols such as propylene glycol, polyethylene glycol, polypropylene glycol, glycol ether, glycerol, and polyoxyethylene alcohols. In certain embodiments herein, particularly in the solution formulations provided herein, the compositions contain a glycol. In other embodiments, the compositions contain propylene glycol and/or polyethylene glycol, including polyethylene glycol

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400. In these embodiments, the glycol is present at a concentration of about 0 mg/mL to about 100 mg/mL, or about 5 mg/mL to about 50 mg/mL. In further embodiments, the compositions contain propylene glycol at a concentration of 17 mg/mL or 20 mg/mL. In other

5 embodiments, the compositions contain polyethylene glycol 400 at a concentration of 10 mg/mL, 20 mg/mL, 30 mg/mL or 50 mg/mL.

The compositions provided herein, particularly the suspension formulations provided herein, may also contain one or more emulsifiers.

10 The particular emulsifier for use in the compositions for long term storage provided herein may be determined empirically using methods well known to those of skill in the art (see, e.g., the Examples). Emulsifiers for use herein include, but are not limited to, polyoxyethylene sorbitan fatty esters or polysorbates, including, but not limited to, polyethylene sorbitan monooleate (Polysorbate 80), polysorbate 20 (polyoxyethylene (20)

15 sorbitan monolaurate), polysorbate 65 (polyoxyethylene (20) sorbitan tristearate), polyoxyethylene (20) sorbitan mono-oleate, polyoxyethylene (20) sorbitan monopalmitate, polyoxyethylene (20) sorbitan monostearate; lecithins; alginic acid; sodium alginate; potassium alginate; ammonium alginate; calcium alginate; propane-1,2-diol alginate; agar;

20 carrageenan; locust bean gum; guar gum; tragacanth; acacia; xanthan gum; karaya gum; pectin; amidated pectin; ammonium phosphatides; microcrystalline cellulose; methylcellulose; hydroxypropylcellulose; hydroxypropylmethylcellulose; ethylmethylcellulose; carboxymethylcellulose; sodium, potassium and calcium salts of fatty

25 acids; mono-and di-glycerides of fatty acids; acetic acid esters of mono-and di-glycerides of fatty acids; lactic acid esters of mono-and di-glycerides of fatty acids; citric acid esters of mono-and di-glycerides of fatty acids; tartaric acid esters of mono-and di-glycerides of fatty acids; mono-and diacetyltartaric acid esters of mono-and di-glycerides of fatty

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acids; mixed acetic and tartaric acid esters of mono-and di-glycerides of fatty acids; sucrose esters of fatty acids; sucroglycerides; polyglycerol esters of fatty acids; polyglycerol esters of polycondensed fatty acids of castor oil; propane-1,2-diol esters of fatty acids; sodium stearyl-2-

5 lactylate; calcium stearyl-2-lactylate; stearyl tartrate; sorbitan monostearate; sorbitan tristearate; sorbitan monolaurate; sorbitan mono-oleate; sorbitan monopalmitate; extract of quillaia; polyglycerol esters of dimerised fatty acids of soya bean oil; oxidatively polymerised soya bean oil; and pectin extract.

10 In certain embodiments herein, the emulsifier(s) is (are) a polyoxyethylene sorbitan fatty ester or polysorbate, including, but not limited to, polyethylene sorbitan monooleate (Polysorbate 80), polysorbate 20 (polyoxyethylene (20) sorbitan monolaurate), polysorbate 65 (polyoxyethylene (20) sorbitan tristearate), polyoxyethylene (20)
15 sorbitan mono-oleate, polyoxyethylene (20) sorbitan monopalmitate, polyoxyethylene (20) sorbitan monostearate; sorbitan monostearate; sorbitan tristearate; sorbitan monolaurate; sorbitan mono-oleate; or sorbitan monopalmitate. In further embodiments, the emulsifier(s) is (are) polysorbate 80, sorbitan monolaurate or polyoxyethylene (20) sorbitan
20 monolaurate.

C. Preparation of compounds for use in the compositions

The preparation of the compounds used in the compositions provided herein is described below. Any such compound or similar compound may be synthesized according to a method discussed in
25 general below or by only minor modification of the methods by selecting appropriate starting materials.

Formoterol may be prepared according to the method disclosed in U.S. Patent No. 3,994,974. Briefly, 4-benzoyloxy-3-nitro- α -bromoacetophenone is reacted with N-benzyl-N-(1-methyl-2-p-

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methoxyphenylethyl)amine to form the α -aminoacetophenone. This compound was subjected to the following series of reactions: (i) reduction of the ketone with sodium borohydride; (ii) reduction of the nitro group with aqueous hydrochloric acid and iron powder; (iii) amine
5 formylation with acetic anhydride and formic acid; and (iv) catalytic reduction over 10% palladium on carbon to afford formoterol free base. Crystallization of the $\frac{1}{2}$ fumarate salt from ethanol provides (formoterol) $\cdot \frac{1}{2}$ fumarate.

The individual enantiomers of formoterol, 2-hydroxy-5-((1S)-1-
10 hydroxy-2-(((1S)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)-formanilide and 2-hydroxy-5-((1R)-1-hydroxy-2-(((1R)-2-(p-methoxyphenyl)-1-methylethyl)amino)ethyl)formanilide, may be prepared by the method disclosed in U.S. Patent No. 6,040,344. Briefly, reaction of optically pure 4-benzyloxy-3-formamidostyrene oxide with an optically
15 pure 4-methoxy- α -methyl-N-(phenylmethyl)benzeneethanamine, followed by debenzylation, affords the desired enantiomer of formoterol. Debenzylation may be accomplished by reduction with hydrogen gas in the presence of a noble metal catalyst, such as palladium on carbon.

The required optically pure 4-benzyloxy-3-formamidostyrene oxide
20 may be prepared from 4-benzyloxy-3-nitro- α -bromoacetophenone by (i) reduction with vorane in the presence of an optically pure aminoindanol, (ii) hydrogenation over platinum oxide catalyst, (iii) formylation with formic acid and acetic anhydride, and (iv) epoxide formation in the presence of potassium carbonate.

25 The required optically pure 4-methoxy- α -methyl-N-(phenylmethyl)-benzeneethanamine may be prepared from 4-methoxyphenylacetone by (i) reductive amination with benzylamine in the presence of hydrogen and a platinum catalyst, and (ii) crystallization of the desired optically pure amine from the resulting racemic mixture as its mandelic acid salt.

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Budesonide may be synthesized by the procedure disclosed in U.S. Patent No. 3,929,768. Briefly, reaction of triamcinolon with propionaldehyde and catalytic perchloric acid in dry dioxane at ambient temperature provides, following chromatography on hydroxy-propylated, cross-linked dextran gel, budesonide.

Fluticasone propionate may be synthesized by the procedure disclosed in U.S. Patent No. 4,335,121. Briefly, the corresponding carbothioic acid is prepared from the carboxylic acid precursor by reaction with dimethylthiocarbamoyl chloride in the presence of triethylamine.

Reaction with bromochloromethane and sodium hydrogen carbonate in dimethylacetamide affords the corresponding S-chloromethyl carbothioate. This compound is treated with sodium iodide in acetone to give the corresponding S-iodomethyl carbothioate. Fluoride substitution of the iodo group is accomplished by reaction with silver fluoride in acetonitrile to afford the desired compound.

D. Formulation of pharmaceutical compositions

The compositions provided herein are prepared by procedures well known to those of skill in the art. For example, a solution formulations may be prepared by the procedure of EXAMPLE 1. Briefly, polyethylene glycol 400 and/or propylene glycol, and a preservative, such as vitamin E TPGS, are mixed at about 42 °C until a homogeneous solution forms. The temperature is lowered and the steroidal anti-inflammatory agent is added. In a second vessel, formoterol fumarate dihydrate and the remaining ingredients are dissolved in approximately 70% water. The two solutions are mixed and the resulting solution is diluted with water to the desired volume.

Suspension formulations are prepared by the procedure of EXAMPLE 2. Briefly, all ingredients other than the steroidal anti-inflammatory agent and formoterol fumarate dihydrate are dissolved in

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about 40% water with mixing. The steroidal anti-inflammatory agent, which is micronized, is dispersed in the above mixture with high speed homogenization. Formoterol fumarate dihydrate is dissolved in 50% water and added to the steroidal suspension with mixing until a uniform

5 suspension forms.

E. Evaluation of the activity of the compositions

Standard physiological, pharmacological and biochemical procedures are available for testing the compositions provided herein to identify those that possess bronchodilatory activity.

- 10 In vitro and in vivo assays that may be used to evaluate bronchodilatory activity are well known to those of skill in the art. See also, e.g., U.S. Patent Nos. 3,994,974, and 6,068,833; German Patent No. 2,305,092; Kaumann et al. (1985) Naunyn-Schmied Arch. Pharmacol. 331:27-39; Lemoine et al. (1985) Naunyn-Schmied Arch. Pharmacol. 331:40-51; Tomioka et al. (1981) Arch. Int. Pharmacodyn. 250:279-292; Dellamary et al. (2000) Pharm. Res. 17(2):168-174; Rico-Mendez et al. (1999) Rev. Alerg. Mex. 46(5):130-135; Seberova et al. (2000) Respir. Med. 94(6):607-611; Lotvall et al. (1999) Can. Respir. J. 6(5):412-416; Campbell et al. (1999) Respir. Med. 93(4):236-244;
- 15 Nightingale et al. (1999) Am. J. Respir. Crit. Care Med. 159(6):1786-1790; Lecaillon et al. (1999) Eur. J. Clin. Pharmacol. 55(2):131-138; Bartow et al. (1998) Drugs 55(2):303-322; Ekstrom et al. (1998) Respir. Med. 92(8):1040-1045; Ringdal et al. (1998) Respir. Med. 92(8):1017-1021; Totterman et al. (1998) Eur. Respir. J. 12(3):573-579; Palmqvist
- 20 et al. (1997) Eur. Respir. J. 10(11):2484-2489; Nielsen et al. (1997) Eur. Respir. J. 10(9):2105-2109; Ullman et al. (1996) Allergy 51(10):745-748; Selroos et al. (1996) Clin. Immunother. 6:273-299; and Schreurs et al. (1996) Eur. Respir. J. 9(8):1678-1683.

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Ingredient	Concentration	
	Low strength	High strength
Citrate buffer	5 mM	5 mM
Sodium Chloride	7.5 mg/mL	6.8 mg/mL
Water	q.s.	q.s.

5 Fluticasone propionate/Formoterol solution formulations

	Ingredient	Concentration	
		Low strength	High strength
10	Formoterol fumarate dihydrate	85 μ g/mL	170 μ g/mL
	Fluticasone propionate	125 μ g/mL	250 μ g/mL
	Vitamin E TPGS	30 mg/mL	50 mg/mL
15	Either: Propylene glycol or Polyethylene glycol 400	17 mg/mL 30 mg/mL	20 mg/mL 50 mg/mL
	Citrate buffer	5 mM	5 mM
20	Sodium Chloride	1.5 mg/mL	0 mg/mL
	Water	q.s.	q.s.

EXAMPLE 2

Preparation of Suspension Formulations

25 All ingredients, with the exception of formoterol fumarate dihydrate and the steroidal anti-inflammatory agent, e.g., budesonide or fluticasone propionate, were dissolved in about 40% water in a container with mixing. The steroidal active ingredient was added and the mixture was

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dispersed with high speed homogenization. Formoterol fumarate dihydrate was dissolved in about 50% water with mixing and the resulting solution was added to the steroidal suspension with mixing until a uniform suspension formed.

- 5 Using the above procedure, the following suspension formulations containing the indicated ingredients in the indicated amounts were prepared.

Budesonide/Formoterol suspension formulations

10

Ingredient	Concentration
Formoterol fumarate dihydrate	5-2000 $\mu\text{g/mL}$ or 50-200 $\mu\text{g/mL}$
Budesonide	125-500 $\mu\text{g/mL}$
Disodium edetate	0.1-0.2 mg/mL
15 Polysorbate 80	0.2-0.3 mg/mL
Sodium chloride	5-10 mg/mL
Citrate buffer	1-20 mM
Water	q.s.

20

Fluticasone propionate/Formoterol suspension formulations

25

Ingredient	Concentration
Formoterol fumarate dihydrate	5-2000 $\mu\text{g/mL}$ or 50-200 $\mu\text{g/mL}$
Fluticasone propionate micronized	250-1000 $\mu\text{g/mL}$
25 Sorbitan monolaurate	0.05-0.2 mg/mL
Polyoxyethylene 20 sorbitan monolaurate	0.1-0.3 mg/mL
Sodium chloride	5-10 mg/mL
Citrate buffer	1-20 mM
30 Water	q.s.

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Since modifications will be apparent to those of skill in this art, it is intended that this invention be limited only by the scope of the appended claims.

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WHAT IS CLAIMED IS:

1. A pharmaceutical composition, comprising (i) formoterol, or a derivative thereof; and (ii) a steroidal anti-inflammatory agent, or a derivative thereof;
- 5 in a pharmacologically suitable fluid, wherein the composition is stable during long term storage and the fluid comprises water.
2. The pharmaceutical composition of claim 1, wherein the composition has an estimated shelf-life of greater than 1 month usage time at 25 °C and greater than or equal to 1 year storage time at 5 °C.
- 10 3. The pharmaceutical composition of claim 1 or 2, wherein greater than about 80% of the initial formoterol is present after 1 month usage time at 25 °C and 1 year storage time at 5 °C.
4. The pharmaceutical composition of any of claims 1-3, wherein the pharmacologically suitable fluid comprises a polar solvent.
- 15 5. The pharmaceutical composition of claim 4, wherein the polar solvent is a protic solvent.
6. The pharmaceutical composition of any of claims 1-5, further comprising a tonicity adjusting agent.
7. The pharmaceutical composition of claim 6, wherein the
- 20 tonicity adjusting agent is ammonium carbonate, ammonium chloride, ammonium lactate, ammonium nitrate, ammonium phosphate, ammonium sulfate, ascorbic acid, bismuth sodium tartrate, boric acid, calcium chloride, calcium disodium edetate, calcium gluconate, calcium lactate, citric acid, dextrose, diethanolamine, dimethylsulfoxide, edetate disodium,
- 25 edetate trisodium monohydrate, fluorescein sodium, fructose, galactose, glycerin, lactic acid, lactose, magnesium chloride, magnesium sulfate, mannitol, polyethylene glycol, potassium acetate, potassium chlorate, potassium chloride, potassium iodide, potassium nitrate, potassium phosphate, potassium sulfate, propylene glycol, silver nitrate, sodium

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acetate, sodium bicarbonate, sodium biphosphate, sodium bisulfite, sodium borate, sodium bromide, sodium cacodylate, sodium carbonate, sodium chloride, sodium citrate, sodium iodide, sodium lactate, sodium metabisulfite, sodium nitrate, sodium nitrite, sodium phosphate, sodium
5 propionate, sodium succinate, sodium sulfate, sodium sulfite, sodium tartrate, sodium thiosulfate, sorbitol, sucrose, tartaric acid, triethanolamine, urea, urethan, uridine or zinc sulfate.

8. The pharmaceutical composition of claim 6 or 7, wherein the tonicity adjusting agent is sodium chloride.

10 9. The pharmaceutical composition of any of claims 1-3, wherein the pharmacologically suitable fluid comprises a buffer.

10. The pharmaceutical composition of any of claims 1-8, further comprising a buffer.

11. The pharmaceutical composition of claim 9 or 10, wherein
15 the buffer is citric acid/phosphate, acetate, barbital, borate, Britton-Robinson, cacodylate, citrate, collidine, formate, maleate, McIlvaine, phosphate, Prideaux-Ward, succinate, citrate-phosphate-borate (Teorell-Stanhagen), veronal acetate, MES (2-(N-morpholino)ethanesulfonic acid), BIS-TRIS (bis(2-hydroxyethyl)iminotris(hydroxymethyl)methane), ADA (N-
20 (2-acetamido)-2-iminodiacetic acid), ACES (N-(carbamoylmethyl)-2-aminoethanesulfonic acid), PIPES (piperazine-N,N'-bis(2-ethanesulfonic acid)), MOPSO (3-(N-morpholino)-2-hydroxypropanesulfonic acid), BIS-TRIS PROPANE (1,3-bis(tris(hydroxymethyl)methylamino)propane), BES (N,N-bis(2-hydroxyethyl)-2-aminoethanesulfonic acid), MOPS (3-(N-
25 morpholino)propanesulfonic acid), TES (N-tris(hydroxymethyl)methyl-2-aminoethanesulfonic acid), HEPES (N-(2-hydroxyethyl)piperazine-N'-(2-ethanesulfonic acid), DIPSO (3-(N,N-bis(2-hydroxyethyl)amino)-2-hydroxypropanesulfonic acid), MOBS (4-(N-morpholino)butanesulfonic acid), TAPSO (3-(N-tris(hydroxymethyl)methylamino)-2-hydroxy-

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propanesulfonic acid), tris(hydroxymethylaminomethane, HEPPSO (N-(2-hydroxyethyl)piperazine-N'-(2-hydroxypropanesulfonic acid), POPSO (piperazine-N,N'-bis(2-hydroxypropanesulfonic acid)), TEA (triethanolamine), EPPS (N-(2-hydroxyethyl)piperazine-N'-(3-propane-sulfonic acid), TRICINE (N-tris(hydroxymethyl)methylglycine), GLY-GLY (glycylglycine), BICINE (N,N-bis(2-hydroxyethyl)glycine), HEPBS (N-(2-hydroxyethyl)piperazine-N'-(4-butanesulfonic acid)), TAPS (N-tris(hydroxymethyl)methyl-3-aminopropanesulfonic acid), or AMPD (2-amino-2-methyl-1,3-propanediol) buffer.

10 12. The pharmaceutical composition of any of claims 9-11, wherein the buffer comprises citric acid/phosphate buffer, acetate buffer, citrate buffer or phosphate buffer.

 13. The pharmaceutical composition of any of claims 9-12, wherein the buffer is citrate buffer.

15 14. The pharmaceutical composition of any of claims 9-13, wherein the buffer concentration is from about 0.01 mM to about 150 mM.

 15. The pharmaceutical composition of any of claims 9-14, wherein the buffer concentration is from about 1 mM to about 50 mM.

20 16. The pharmaceutical composition of any of claims 9-15, wherein the buffer concentration is from about 1 mM to about 20 mM.

 17. The pharmaceutical composition of any of claims 9-16, wherein the buffer concentration is about 20 mM.

 18. The pharmaceutical composition of any of claims 9-16,
25 wherein the buffer concentration is about 5 mM.

 19. The pharmaceutical composition of any of claims 1-18, wherein the ionic strength of the composition is about 0 to about 0.4.

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20. The pharmaceutical composition of any of claims 1-19, wherein the ionic strength of the composition is about 0.05 to about 0.16.

21. The pharmaceutical composition of any of claims 1-20,
5 wherein the pH of the composition is about 2.0 to about 8.0.

22. The pharmaceutical composition of any of claims 1-21, wherein the pH of the composition is about 4.0 to about 6.0.

23. The pharmaceutical composition of any of claims 1-22, wherein the pH of the composition is about 4.5 to about 5.5.

10 24. The pharmaceutical composition of any of claims 1-23, wherein the pH of the composition is about 5.0.

25. The pharmaceutical composition of any of claims 1-24, wherein the formoterol free base concentration is about 5 $\mu\text{g/mL}$ to about 2 mg/mL.

15 26. The pharmaceutical composition of any of claims 1-25, wherein the formoterol free base concentration is about 10 $\mu\text{g/mL}$ to about 1 mg/mL.

27. The pharmaceutical composition of any of claims 1-26, wherein the formoterol free base concentration is about 50 $\mu\text{g/mL}$ to
20 about 200 $\mu\text{g/mL}$.

28. The pharmaceutical composition of any of claims 1-27, wherein the formoterol free base concentration is about 59 $\mu\text{g/mL}$.

29. The pharmaceutical composition of any of claims 1-27, wherein the formoterol free base concentration is about 118 $\mu\text{g/mL}$.

25 30. The pharmaceutical composition of any of claims 9-29, wherein the buffer is citrate buffer; the buffer concentration is about 5 mM; the ionic strength of the composition is about 0.05 to about 0.16; and the pH of the composition is about 5.0.

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31. The pharmaceutical composition of any of claims 1-30,
wherein the steroidal anti-inflammatory agent is beclomethasone
dipropionate (BDP), beclomethasone monopropionate (BMP), flunisolide,
triamcinolone acetonide, dexamethasone, tipredane, ciclesonid,
5 rofleponide, mometasone, mometasone furoate, RPR 106541, fluticasone
or fluticasone propionate or budesonide, or a derivative thereof.

32. The pharmaceutical composition of any of claims 1-31,
wherein the steroidal anti-inflammatory agent is budesonide or fluticasone
propionate, or a derivative thereof.

10 33. The pharmaceutical composition of any of claims 1-32,
wherein the steroidal anti-inflammatory agent is budesonide, or a
derivative thereof.

34. The pharmaceutical composition of any of claims 31-33,
wherein the budesonide concentration is about 5 $\mu\text{g/mL}$ to about 2
15 mg/mL .

35. The pharmaceutical composition of any of claims 31-34,
wherein the budesonide concentration is about 75 $\mu\text{g/mL}$ to about 500
 $\mu\text{g/mL}$.

36. The pharmaceutical composition of any of claims 31-35,
20 wherein the budesonide concentration is about 125 $\mu\text{g/mL}$ or about 250
 $\mu\text{g/mL}$.

37. The pharmaceutical composition of any of claims 1-32,
wherein the steroidal anti-inflammatory agent is fluticasone propionate.

38. The pharmaceutical composition of any of claims 31, 32 or
25 37, wherein the concentration of fluticasone propionate is about 5 $\mu\text{g/mL}$
to about 2 mg/mL .

39. The pharmaceutical composition of any of claims 31, 32, 37
or 38, wherein the concentration of fluticasone propionate is about 75
 $\mu\text{g/mL}$ to about 1000 $\mu\text{g/mL}$.

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40. The pharmaceutical composition of any of claims 31, 32 or 37-39, wherein the concentration of fluticasone propionate is about 125 $\mu\text{g/mL}$ or about 250 $\mu\text{g/mL}$.

41. The pharmaceutical composition of any of claims 1-40,
5 further comprising one or more of (a) to (j) as follows: (a) a β_2 -adrenoreceptor agonist; (b) a dopamine (D_2) receptor agonist; (c) an IL-5 inhibitor; (d) an antisense modulator of IL-5; (e) a tryptase inhibitor; (f) a tachykinin receptor antagonist; (g) milrinone or milrinone lactate; (h) a leukotriene receptor antagonist; (i) a 5-lipoxygenase inhibitor; or (j) an
10 anti-IgE antibody.

42. The pharmaceutical composition of any of claims 9-29, wherein the buffer is citrate buffer; the buffer concentration is about 20 mM; the ionic strength of the composition is about 0.05 to about 0.16; and the pH of the composition is about 5.0.

15 43. The pharmaceutical composition of any of claims 1-42, further comprising an anticholinergic agent.

44. The pharmaceutical composition of claim 43, wherein the anticholinergic agent is ipratropium bromide, oxitropium bromide, atropine methyl nitrate, tiotropium bromide or glycopyrronium bromide.

20 45. The pharmaceutical composition of claim 43 or 44, wherein the anticholinergic agent is ipratropium bromide.

46. The pharmaceutical composition of claim 45, wherein the ipratropium bromide is present at a concentration of about 5 $\mu\text{g/mL}$ to about 5 mg/mL.

25 47. The pharmaceutical composition of claim 43 or 44, wherein the anticholinergic agent is tiotropium bromide.

48. The pharmaceutical composition of claim 47, wherein the tiotropium bromide is present at a concentration of about 5 $\mu\text{g/mL}$ to about 5 mg/mL.

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49. The pharmaceutical composition of any of claims 1-48 that is a solution.

50. The pharmaceutical composition of any of claims 1-50 that is a suspension.

5 51. The pharmaceutical composition of any of claims 1-50 that has been nebulized.

52. A nebulized suspension or solution, comprising (i) formoterol or a derivative thereof; and (ii) a steroidal anti-inflammatory agent, or a derivative thereof; in a pharmacologically suitable fluid.

10 53. A kit, comprising:

(a) an aqueous composition comprising (i) formoterol or a derivative thereof, and (ii) a steroidal anti-inflammatory agent or a derivative thereof, formulated for single dosage administration; and

15 (b) a nebulizer.

54. The kit of claim 53, wherein the aqueous composition comprises (a) formoterol free base at a concentration of about 59 $\mu\text{g/mL}$; (b) aqueous saline comprising sodium chloride; and (c) citrate buffer at a concentration of about 5 mM;

20 wherein the ionic strength of the composition is about 0.05 to about 0.16; and the pH of the composition is about 5.0.

55. The kit of claim 53, wherein the aqueous composition comprises (a) formoterol free base at a concentration of about 118 $\mu\text{g/mL}$; (b) aqueous saline comprising sodium chloride; and (c) citrate
25 buffer at a concentration of about 5 mM;

wherein the ionic strength of the composition is about 0.05 to about 0.16; and the pH of the composition is about 5.0.

56. A kit, comprising:

(a) the pharmaceutical composition of any of claims 1-50; and

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(b) a nebulizer.

57. The kit of claim 53, wherein the aqueous composition comprises (a) formoterol free base at a concentration of about 59 $\mu\text{g/mL}$; (b) aqueous saline comprising sodium chloride; and (c) citrate buffer at a
5 concentration of about 20 mM;

wherein the ionic strength of the composition is about 0.05 to about 0.16; and the pH of the composition is about 5.0.

58. The kit of claim 53, wherein the aqueous composition comprises (a) formoterol free base at a concentration of about 118
10 $\mu\text{g/mL}$; (b) aqueous saline comprising sodium chloride; and (c) citrate buffer at a concentration of about 20 mM;

wherein the ionic strength of the composition is about 0.05 to about 0.16; and the pH of the composition is about 5.0.

59. A combination, comprising:

15 (a) the pharmaceutical composition of any of claims 1-50 formulated for single dosage administration; and
(b) a vial.

60. The combination of claim 59, wherein the aqueous composition comprises (a) formoterol free base at a concentration of
20 about 59 $\mu\text{g/mL}$; (b) aqueous saline comprising sodium chloride; and (c) citrate buffer at a concentration of about 5 mM;

wherein the ionic strength of the composition is about 0.05 to about 0.16; and the pH of the composition is about 5.0.

61. The combination of claim 59, wherein the aqueous
25 composition comprises (a) formoterol free base at a concentration of about 118 $\mu\text{g/mL}$; (b) aqueous saline comprising sodium chloride; and (c) citrate buffer at a concentration of about 5 mM;

wherein the ionic strength of the composition is about 0.05 to about 0.16; and the pH of the composition is about 5.0.

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62. The combination of claim 59, wherein the aqueous composition comprises (a) formoterol free base at a concentration of about 59 $\mu\text{g/mL}$; (b) aqueous saline comprising sodium chloride; and (c) citrate buffer at a concentration of about 20 mM;

5 wherein the ionic strength of the composition is about 0.05 to about 0.16; and the pH of the composition is about 5.0.

63. The combination of claim 59, wherein the aqueous composition comprises (a) formoterol free base at a concentration of about 118 $\mu\text{g/mL}$; (b) aqueous saline comprising sodium chloride; and (c)
10 citrate buffer at a concentration of about 20 mM;

wherein the ionic strength of the composition is about 0.05 to about 0.16; and the pH of the composition is about 5.0.

64. A combination, comprising:

a composition comprising formoterol, or a derivative thereof, in a
15 pharmacologically suitable fluid, wherein the composition is stable during long term storage and the fluid comprises water; and

a composition comprising a bronchodilating steroid, or a derivative thereof.

65. The combination of claim 64, further comprising a nebulizer.

20 66. The combination of claim 65 that is packaged as a kit; optionally comprising instructions for use of the nebulizer; and optionally comprising instructions for mixing the compositions.

67. A method for the treatment, prevention, or amelioration of one or more symptoms of bronchoconstrictive disorders, comprising
25 administering an effective amount of the pharmaceutical composition of any of claims 1-50 to a subject in need of such treatment.

68. The method of claim 67, further comprising administering one or more of (a) to (j) as follows: (a) a β_2 -adrenoreceptor agonist; (b) a dopamine (D_2) receptor agonist; (c) an IL-5 inhibitor; (d) an antisense

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modulator of IL-5; (e) a tryptase inhibitor; (f) a tachykinin receptor antagonist; (g) milrinone or milrinone lactate; (h) a leukotriene receptor antagonist; (i) a 5-lipoxygenase inhibitor; or (j) an anti-IgE antibody; simultaneously with, prior to or subsequent to the formoterol/steroidal
5 anti-inflammatory agent composition.

69. A method for the treatment, prevention, or amelioration of one or more symptoms of bronchoconstrictive disorders, comprising:

(i) administering an effective amount of a pharmaceutical composition comprising formoterol, or a derivative thereof, in a
10 pharmacologically suitable fluid, wherein the composition is stable during long term storage and the fluid comprises water; and

(ii) simultaneously with, prior to, or subsequent to administering the formoterol composition, administering an effective amount of a pharmaceutical composition comprising a steroidal anti-inflammatory
15 agent in a pharmacologically acceptable carrier.

70. The method of claim 68, wherein the steroidal anti-inflammatory agent is beclomethasone dipropionate (BDP), beclomethasone monopropionate (BMP), flunisolide, triamcinolone acetonide, dexamethasone, tipredane, ciclesonid, rofleponide,
20 mometasone, mometasone furoate, RPR 106541, fluticasone or fluticasone propionate or budesonide, or a derivative thereof.

71. The method of claim 70, wherein the steroidal anti-inflammatory agent is fluticasone propionate or budesonide, or a derivative thereof.

25 72. The method of claim 70 or 71, wherein the steroidal anti-inflammatory agent is budesonide, or a derivative thereof.

73. The method of claim 70 or 71, wherein the steroidal anti-inflammatory agent is fluticasone propionate, or a derivative thereof.

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74. An article of manufacture, comprising packaging material, an aqueous composition comprising the composition of any of claims 1-50 formulated for single dosage administration, which is useful for treatment, prevention or amelioration of one or more symptoms of
- 5 diseases or disorders associated with undesired and/or uncontrolled bronchoconstriction, and a label that indicates that the composition is used for treatment, prevention or amelioration of one or more symptoms of diseases or disorders associated with undesired and/or uncontrolled bronchoconstriction.